## We Claim:

- A particulate complex comprising a nucleic acid and a biodegradable cationized polyhydroxylated molecule, wherein said molecule has a charge up to approximately 1.0 meq/g.
- A complex according to claim 1, wherein the nucleic acid is double or single stranded DNA or RNA, or a mixture thereof.
- A complex according to claim 1, wherein the nucleic acid is a natural or chemically modified oligonucleotide or a derivative thereof.
- A complex according to claim 1, wherein the nucleic acid is a natural or chemically modified polynucleotide or a derivative thereof.
- A complex according to claim 1, wherein the biodegradable cationized polyhydroxylated molecule has a charge between approximately 0.1 and approximately 0.85 meq/g.
- A complex according to claim 1, wherein the polyhydroxylated molecule is a saccharide comprising a cationic moiety.
- 7. A complex according to claim 6, wherein the saccharide is a polysaccharide
- 8. A complex according to claim 6, wherein the saccharide is an oligosaccharide.
- 9. A complex according to claim 6, wherein the saccharide is a monosaccharide.
- 10. A complex according to claim 6, wherein the cationic moiety comprises a secondary or tertiary amino group; quaternary ammonium ion; or a combination thereof.

- A complex according to claim 10, wherein the quaternary ammonium ion is glycidyl trimethylammonium.
- A complex according to claim 1, wherein the cationized polyhydroxylated molecule has a molecular weight of between approximately 0.18 kDa and approximately 1000 kDa.
- A complex according to claim 12, wherein the cationized polyhydroxylated molecule has a molecular weight of between about 0.5kDa and 500kDa.
- 14. A complex according to claim 1, wherein the complex is of size between approximately 100 nm to approximately 10  $\mu$ m.
- 15. A complex according to claim 1, wherein the complex has a charge ratio of cationized polyhydroxylated molecule to nucleic acid between approximately 0.3 to 1, and wherein the complex is globally negative.
- 16. A complex according to claim 1, wherein the complex has a charge ratio of cationized polyhydroxylated molecule to nucleic acid between 1 to approximately 20, and wherein the complex is globally positive.
- 17. A solution comprising a complex according to claim 16, wherein the solution further comprises excess cationized polyhydroxylated molecule that is not complexed to the nucleic acid.
- 18. A method for protecting a nucleic acid molecule when transfecting said molecule into a cell, said method comprising complexing the nucleic acid with a cationized polyhydroxylated molecule to form a particulate complex according to claim 1.

- A method according to claim 18, wherein the complex has a charge ratio of cationized polyhydroxylated molecule to nucleic acid between approximately 0.3 and approximately 20.
- 20. A method for transfecting a nucleic acid molecule into a cell ex vivo, said method comprising complexing the nucleic acid with a cationized polyhydroxylated molecule to form a particulate complex according to claim 1, and transfecting the cell with the complex.
- A method according to claim 20, wherein the complex has a charge ratio of cationized polyhydroxylated molecule to nucleic acid between approximately 0.3 and approximately 20.
- 22. A method for administering a nucleic acid molecule to a mammal, said method comprising complexing the nucleic acid with a cationized polyhydroxylated molecule to form a particulate complex according to claim 1, and administering the complex to the mammal.
- 23. A method according to claim 22, wherein the complex has a charge ratio of cationized polyhydroxylated molecule to nucleic acid between approximately 0.3 and approximately 20.
- 24. A method according to claim 22 wherein the administration of the complex is intramuscular
- A method according to claim 22, wherein the nucleic acid encodes an immunogenic antigen.
- A method according to claim 22, wherein the nucleic acid encodes a therapeutic protein.
- 27. A pharmaceutical composition comprising the complex of claim 1.

- 28. A pharmaceutical composition according to claim 27 further comprising a transfection enhancer.
- 29. A pharmaceutical composition according to claim 28, wherein said transfection enhancer is selected from the group consisting of lipids, detergents, enzymes, peptides, and enzyme inhibitors.
- 30. A pharmaceutical composition according to claim 28, wherein said transfection enhancer comprises free cationized polyhydroxylated molecules not complexed to the nucleic acid.